

*Supporting Information Available*

## Antibacterial and photochemical properties of cellulose nanofibers–titania nanocomposites loaded with two different types of antibiotic medicines

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### Surface modification

Cellulose nanofibers surface were modified by Phosphomycin and Tetracycline in amounts calculated to achieve uniform, single layer coverage. The surface of titania was estimated using simple mathematical model where the particles are spherical, uniform and homogenous. According to our assumption that drug bonded with the biopolymer through interaction with TiO<sub>2</sub>, we firstly calculated the total number of titania nanoparticles, which was added to the cellulose nanofibers, by following formula:

$$N_{total} = m_{total}/m_p$$

where  $m_{total}$  – total mass of TiO<sub>2</sub> particles (g),  $m_p$  – mass of the single TiO<sub>2</sub> particle (g).

The total surface area of titania nanoparticles grafted onto cellulose nanofibers can be estimated as follows:

$$S_{total} = N_{total} \cdot S_p$$

where  $N_{total}$  – total number of titania nanoparticles,  $S_p$  - surface of the single TiO<sub>2</sub> particle (nm<sup>2</sup>). Then, the amount of drug  $n$  (mol) can be found as follows:

$$n = S_{total} \cdot N_A / S$$

where  $S$  – area of functional group (nm<sup>2</sup>) and  $N_A$  is Avogadro number. The approximate area of 0.24 nm<sup>2</sup> per phosphonate group is commonly proposed in literature<sup>1</sup> and utilized by us in our previous papers<sup>2,3</sup>.

**Table S1** Antimicrobial activity of samples against *S.aureus* and *E.coli*. The means (mm) ± standard deviation for at least three replicates are illustrated. (\*na = no activity)

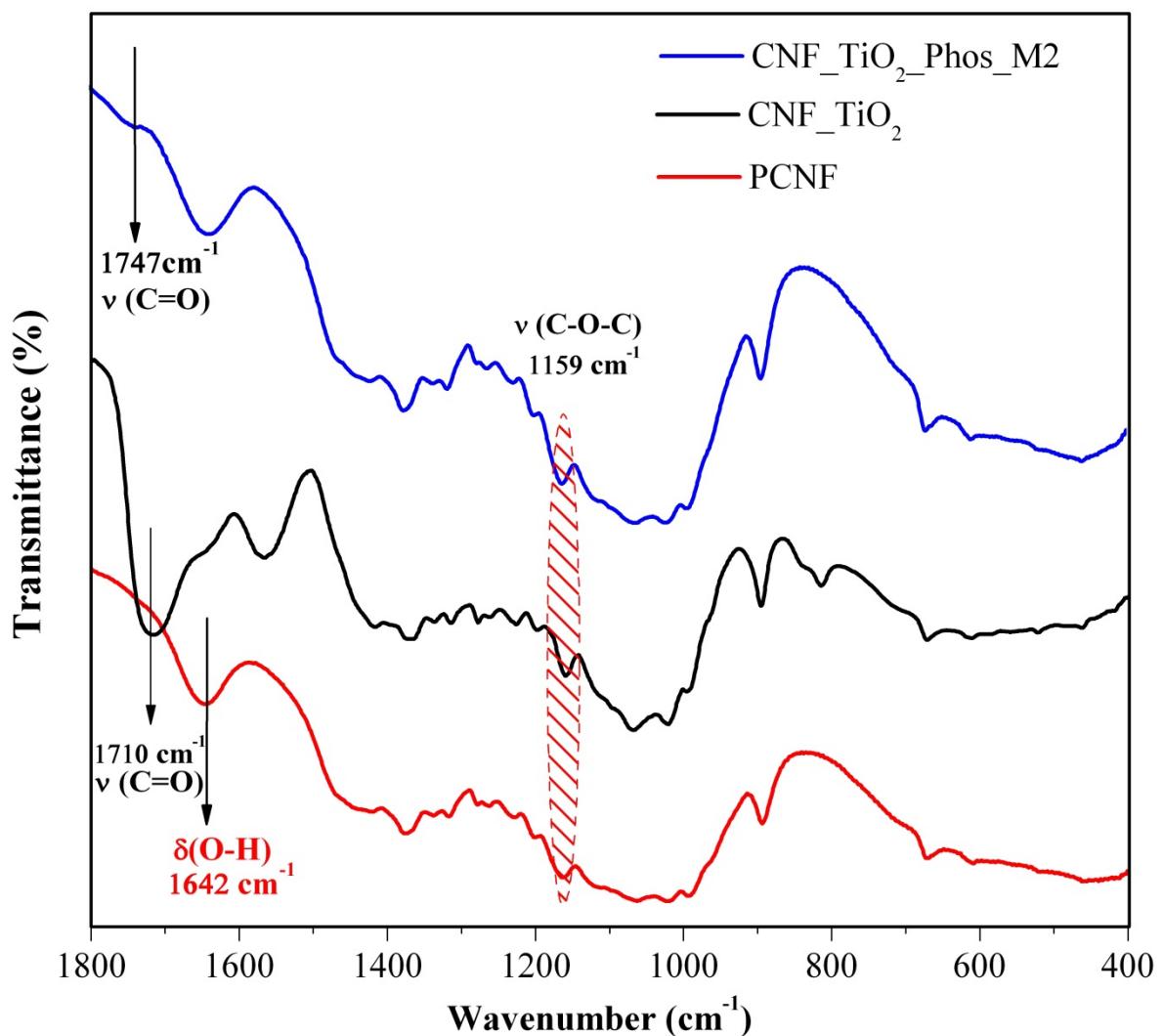
Sample	<i>S.aureus</i>		<i>E.coli</i>	
	Concentration (µg/disk)	Inhibition zone (mm)	Concentration (µg/disk)	Inhibition zone (mm)
CNF_TC_M0	1	23.5±2.9	1	14.5±1.7
	10	30.5±0.6	10	22.0±0.8
	50	34.0±1.8	50	27.3±1.7
	100	38.3±2.6	100	32.8±2.2
CNF_TiO <sub>2</sub> _TC_M1	1	19.3±3.4	1	9.3±2.1
	10	25.8±3.7	10	16.3±0.5
	50	32.5±4.7	50	21.0±1.4
	100	32.3±4.4	100	23.3±0.5
CNF_Phos_M0	10	28.5±3.2	50	25.0±0.0
	50	42.0±7.2		
	100	45.8±5.5	100	29.0±2.8
CNF_TiO <sub>2</sub> _Phos_M2	10	na*	50	na
	50	na		
	100	na	100	na

**Table S2** The antimicrobial effect of samples against *S.aureus* obtained from the liquid broth assay

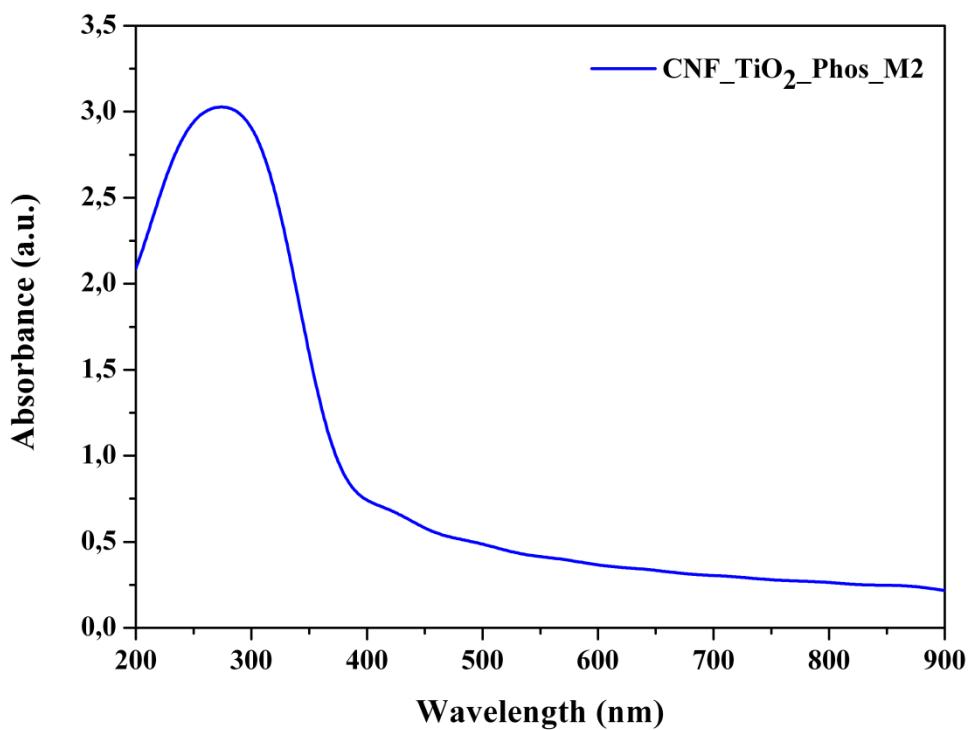
Sample	Concentration (µg/ml)	Time delay (h)	Growth inhibition (%)
CNF_TC_M0	1	0	100
CNF_TiO <sub>2</sub> _TC_M1	1	0	100
CNF_Phos_M0	100	0	100
CNF_TiO <sub>2</sub> _Phos_M2	100	0*	100
		0	96.6
		12	99.2
		18	98.1
		24	96.6

(\* ) with addition of citrate buffer (pH=6,0.03M)

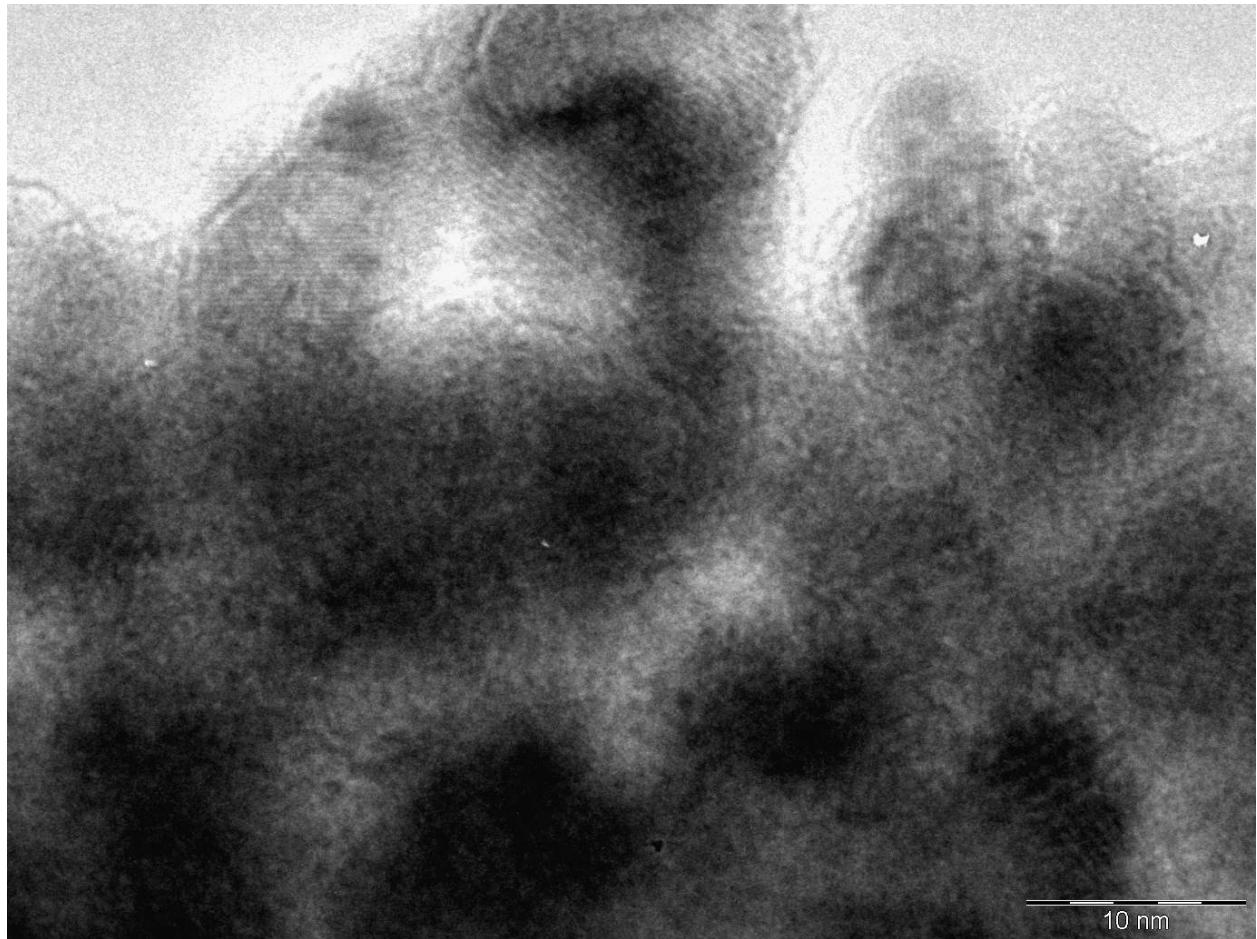
**Figure FS1** FTIR of the CNF\_TiO<sub>2</sub>\_Phos\_M2 compared to CNF\_TiO<sub>2</sub> and PCNF.



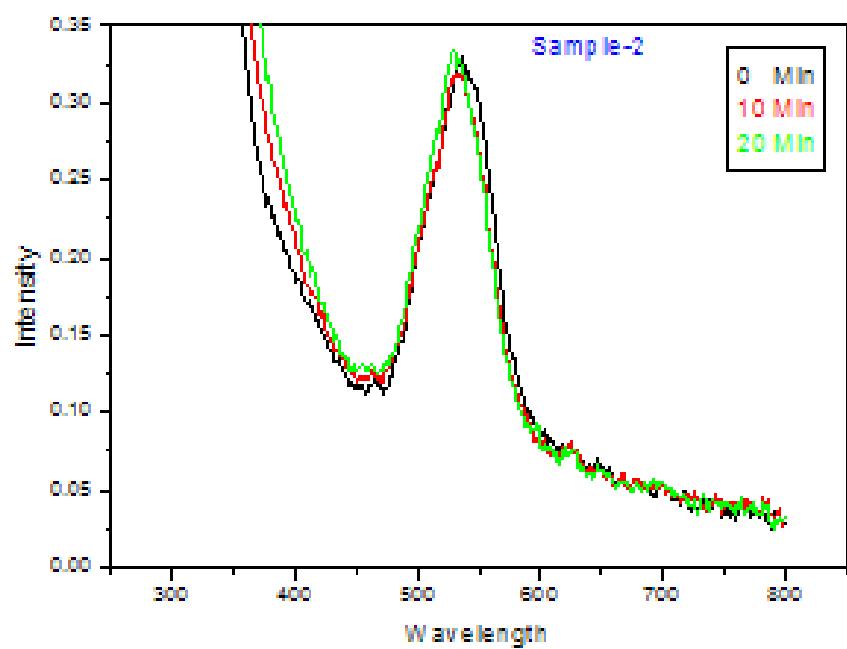
**Figure FS2** UV-Vis spectrum of the CNF\_TiO<sub>2</sub>\_Phos\_M2 material in the form of thin transparent film.



**Figure FS3** High resolution TEM view of the applied TiO<sub>2</sub> nanoparticles (reprinted with permission from Angew. Chem. Int. Ed. 2008, 47, 8506 –8509)



**Figure FS4** The stability demonstration for solutions of Rhodamine B in the presence of the applied TiO<sub>2</sub> nanoparticles (reprinted with permission from the supplementary materials to Angew. Chem. Int. Ed. 2008, 47, 8506 –8509)



## **References**

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